

## METHODS

This invention relates to polymorphisms in the human pyruvate dehydrogenase E1 $\beta$  (PDH E1 $\beta$ ) gene. The invention also relates to methods and materials for analysing allelic  
5 variation in the PDH E1 $\beta$  gene, and to the use of PDH E1 $\beta$  polymorphism in the diagnosis and treatment of diseases in which modulation of pyruvate dehydrogenase activity could be of therapeutic benefit, such as diabetes, asthma, obesity, sepsis and peripheral vascular disease.

The production of energy for biosynthesis of complex molecules and for muscle contraction is mediated by the hydrolysis of high energy phosphate bonds within adenosine  
10 triphosphate (ATP). In oxidative metabolism ATP is generated from acetyl coenzyme A (acetyl CoA), which itself is produced by the beta-oxidation of fatty acids, or as a result of the metabolism of glucose via the glycolytic pathway. The key regulatory enzyme which controls the rate of acetyl CoA formation from glucose is pyruvate dehydrogenase (PDH), which catalyses the oxidation of pyruvate to acetyl CoA and carbon dioxide with concomitant  
15 reduction of NAD to NADH.

PDH is a multienzyme complex located in the mitochondrial matrix, containing multiple copies of three enzyme components required to complete the conversion of pyruvate to acetyl CoA (Patel and Roche 1990; FASEB J., 4: 3224-3233). E1 (pyruvate decarboxylase, E.C. 1.2.4.1) catalyses the non-reversible removal of carbon dioxide from pyruvate; E2  
20 (dihydrolipoamide acetyltransferase, E.C. 2.3.1.12) catalyses the formation of acetyl CoA; and E3 (dihydrolipoamide dehydrogenase, E.C. 1.8.1.4) reduces NAD to NADH. The E1 enzyme is a heterotetramer composed of two  $\alpha$  and two  $\beta$  subunits. Decarboxylation of pyruvate, catalysed by E1 is the rate limiting step in the overall activity of the PDH complex. This step is also the target for a cycle of phosphorylation and dephosphorylation which forms  
25 one of the main mechanisms for regulating PDH activity. Two additional enzyme activities are also associated with the PDH complex: a specific kinase (PDK) which is capable of phosphorylating E1 $\alpha$  at three serine residues, and a loosely-associated specific phosphatase which reverses the phosphorylation. Phosphorylation of only one of the three serine residues on E1 $\alpha$  renders E1 inactive. Removal of the phosphate groups by the specific phosphatase  
30 restores activity. Thus, the proportion of PDH in its active (dephosphorylated) state is determined by a balance between the activity of the kinase and phosphatase. The activity of

the kinase may be regulated in vivo by the relative concentrations of metabolic substrates such as NAD/NADH, CoA/acetylCoA and ADP/ATP as well as by the availability of pyruvate itself, therefore providing highly regulated, responsive control of PDH activity.

Genetic abnormalities in the PDH complex are the most common cause of primary lactic acidosis in humans. The majority of cases have been linked with a defect in the E1 $\alpha$  subunit. Pathologies associated with defects in the PDH complex conform to a broad clinical spectrum ranging from fatal lactic acidosis in the newborn, to a range of chronic neurodegenerative conditions with gross structural abnormalities in the central nervous system. E1 $\alpha$  deficiency is an X-linked disorder which manifests different patterns of clinical presentation between males and females. In addition, heterozygous females show a wide variation in clinical severity of the disease, due largely to variations in the pattern of X-inactivation and differential effects of specific gene mutations on the expression, stability and activity of the mutant protein. A number of mutations in the PDH gene which lead to pyruvate dehydrogenase deficiency have been documented (for a review see NIH OMIM database, reference 312170).

In disease states such as both non-insulin dependent (NIDDM) and insulin-dependent diabetes (IDDM), oxidation of lipids is increased with a concomitant reduction in utilisation of glucose, contributing to the hyperglycaemia. The activity of PDH is reduced in both insulin-dependent and non insulin-dependent diabetes. A further consequence of reduced PDH activity is an increase in pyruvate concentration resulting in increased availability of lactate as a substrate for hepatic gluconeogenesis. Diabetes is further exacerbated by impaired insulin secretion, which has been shown to be associated with reduced PDH activity in pancreatic  $\beta$ -cells. It is believed that increasing the activity of PDH may increase the rate of glucose oxidation and hence overall glucose utilisation, in addition to reducing hepatic glucose output.

Oxidation of glucose is capable of yielding more molecules of ATP per mole of oxygen than is oxidation of fatty acids, therefore in conditions where energy demand may exceed energy supply, such as myocardial ischaemia and reperfusion, intermittent claudication, cerebral ischaemia and reperfusion, shifting the balance of substrate utilisation in favour of glucose metabolism may be expected to improve the ability to maintain ATP levels and hence function. Activation of PDH is predicted to have this effect.

An agent which is capable of activating PDH is expected to be of benefit in treating conditions where an excess of circulating lactic acid is manifest such as in certain cases of sepsis.

The agent dichloroacetic acid which increases the activity of PDH after acute administration in animals (Vary et al., 1988; Circ. Shock, 24: 3-18) has been shown to have the predicted effects in reducing glycaemia (Stacpoole et al, 1978 N. Engl. J. Med. 298, 526-530) and as a therapy for myocardial ischaemia (Bersin and Stacpoole 1997; American Heart Journal, 134: 841-855) and lactic acidemia (Stacpoole et al, 1983 N. Engl. J. Med 309, 390-396).

cDNA sequences encoding PDH E1 $\beta$  have been submitted to public databases under the following accession numbers: X57778, M34479, J03576, M34055, M34056, M54788. A genomic DNA sequence encoding PDH E1 $\beta$  has been submitted under accession number D90086. All positions in the coding region and 3' untranslated region of the human PDH E1 $\beta$  gene herein refer to the positions in SEQ ID NO: 1 (which is equivalent to EMBL accession number X57778 at the time of filing this application) unless stated otherwise or apparent from the context.

The cytogenetic location of the human PDH E1 $\beta$  gene has been mapped to 3p13-q23 (Olsen *et al.*, Am. J. Hum. Genet. 46, 340-349, 1990).

DNA polymorphisms may lead to variations in amino acid sequence and consequently to altered protein structure and functional activity. Polymorphisms may also affect mRNA synthesis, maturation, transportation and stability. Polymorphisms which do not result in amino acid changes (silent polymorphisms) or which do not alter any known consensus sequences may nevertheless have a biological effect, for example by altering mRNA folding or stability.

Knowledge of polymorphisms may be used to help identify patients most suited to therapy with particular pharmaceutical agents (this is often termed "pharmacogenetics"). Pharmacogenetics can also be used in pharmaceutical research to assist the drug selection process. Polymorphisms may be used in mapping the human genome and to elucidate the genetic component of diseases. The reader is directed to the following references for background details on pharmacogenetics and other uses of polymorphism detection: Linder *et al.* (1997), Clinical Chemistry, 43, 254; Marshall (1997), Nature Biotechnology, 15, 1249;

International Patent Application WO 97/40462, Spectra Biomedical; and Schafer *et al.* (1998), Nature Biotechnology, 16, 33.

Clinical trials have shown that patient response to treatment with pharmaceuticals is often heterogeneous. Thus there is a need for improved approaches to pharmaceutical agent  
5 design and therapy.

The present invention is based on the discovery of a single nucleotide polymorphism (SNP) in the coding region of the human PDH E1 $\beta$  gene, and three single nucleotide polymorphisms in the 3' untranslated region (3'UTR) of the human PDH E1 $\beta$  gene.

According to one aspect of the present invention there is provided a method for the  
10 diagnosis of a polymorphism in a PDH E1 $\beta$  gene in a human, which method comprises determining the sequence of the nucleic acid of the human at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1; and determining the status of the human by reference to polymorphism in the PDH E1 $\beta$  gene.

The term human includes both a human having or suspected of having a PDH-  
15 mediated disease and an asymptomatic human who may be tested for predisposition or susceptibility to such disease. At each position the human may be homozygous for an allele or the human may be a heterozygote.

The term 'PDH-mediated disease' means any disease in which changing the level of PDH or changing the activity of PDH would be of therapeutic benefit.

20 The term 'PDH drug' means any drug which changes the level of PDH or changes the activity of PDH. A drug which increases the activity of PDH is preferred.

The term polymorphism includes single nucleotide substitution, nucleotide insertion and nucleotide deletion, which in the case of insertion and deletion includes insertion or deletion of one or more nucleotides at a position of a gene and variable numbers of a repeated  
25 DNA sequence.

Preferably the polymorphisms are further defined as:

Position	Polymorphism	Region
457	A/G	Coding, silent Gly
1191	A/C	3'UTR
1198	C/T	3'UTR
1342	C/A	3'UTR

A preferred method comprises diagnosis of any one of the following haplotypes:

- (a) 1191C 1198C 1342A;
- 5 (b) 1191A 1198C 1342C; or
- (c) 1191C 1198T 1342A.

In one embodiment of the invention preferably the method for diagnosis described herein is one in which the single nucleotide polymorphism at position 457 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 is presence of A and/or G.

- 10 In another embodiment of the invention preferably the method for diagnosis described herein is one in which the single nucleotide polymorphism at position 1191 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 is presence of A and/or C.

- In another embodiment of the invention preferably the method for diagnosis described herein is one in which the single nucleotide polymorphism at position 1198 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 is presence of C and/or T.
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In another embodiment of the invention preferably the method for diagnosis described herein is one in which the single nucleotide polymorphism at position 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 is presence of C and/or A.

- The method for diagnosis is preferably one in which the sequence is determined by a method selected from amplification refractory mutation system and restriction fragment length polymorphism.
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In another aspect of the invention we provide a method for the diagnosis of PDH-mediated disease, which method comprises:

- i) obtaining sample nucleic acid from an individual,

ii) detecting the presence or absence of a variant nucleotide at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1; and

iii) determining the status of the individual by reference to polymorphism in the PDH E1 $\beta$  gene.

Allelic variation at position 457 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 consists of a single base substitution from A (the published base), preferably to G. Allelic variation at position 1191 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 consists of a single base substitution from A (the published base), preferably to C.

Allelic variation at position 1198 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 consists of a single base substitution from C (the published base), preferably to T. Allelic variation at position 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 consists of a single base substitution from C (the published base), preferably to A. The status of the individual may be determined by reference to allelic variation at any one, two, three or all four positions optionally in combination with any other polymorphism in the gene that is (or becomes) known.

The test sample of nucleic acid is conveniently present in a sample of blood, sputum, skin, bronchoalveolar lavage fluid, or other body fluid or tissue obtained from an individual. It will be appreciated that the test sample may equally comprise a nucleic acid sequence corresponding to the sequence in the test sample, that is to say that all or a part of the region in the sample nucleic acid may firstly be amplified using any convenient technique e.g. PCR, before analysis of allelic variation.

It will be apparent to the person skilled in the art that there are a large number of analytical procedures which may be used to detect the presence or absence of variant nucleotides at one or more polymorphic positions of the invention. In general, the detection of allelic variation requires a mutation discrimination technique, optionally an amplification reaction and optionally a signal generation system. Table 1 lists a number of mutation detection techniques, some based on PCR. These may be used in combination with a number of signal generation systems, a selection of which is listed in Table 2. Further amplification techniques are listed in Table 3. Many current methods for the detection of allelic variation are reviewed by Nollau *et al.*, Clin. Chem. 43, 1114-1120, 1997; and in standard textbooks,

for example "Laboratory Protocols for Mutation Detection", Ed. by U. Landegren, Oxford University Press, 1996 and "PCR", 2<sup>nd</sup> Edition by Newton & Graham, BIOS Scientific Publishers Limited, 1997.

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**Abbreviations:**

ALEX™	Amplification refractory mutation system linear extension
APEX	Arrayed primer extension
ARMS™	Amplification refractory mutation system
b-DNA	Branched DNA
CMC	Chemical mismatch cleavage
bp	base pair
COPS	Competitive oligonucleotide priming system
DGGE	Denaturing gradient gel electrophoresis
FRET	Fluorescence resonance energy transfer
IDDM	Insulin-dependent diabetes mellitus
LCR	Ligase chain reaction
MASDA	Multiple allele specific diagnostic assay
NASBA	Nucleic acid sequence based amplification
NIDDM	non-insulin dependent diabetes mellitus
OLA	Oligonucleotide ligation assay
PCR	Polymerase chain reaction
PDH	Pyruvate Dehydrogenase
PDK	Pyruvate Dehydrogenase Kinase
PDK2	Pyruvate Dehydrogenase Kinase Isoenzyme 2
PTT	Protein truncation test
RFLP	Restriction fragment length polymorphism
SDA	Strand displacement amplification
SNP	Single nucleotide polymorphism
SSCP	Single-strand conformation polymorphism analysis
SSR	Self sustained replication
TGGE	Temperature gradient gel electrophoresis
3'UTR	3' Untranslated Region

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Table 1 - Mutation Detection Techniques

**General:** DNA sequencing, Sequencing by hybridisation

**Scanning:** PTT\*, SSCP, DGGE, TGGE, Cleavase, Heteroduplex analysis, CMC, Enzymatic mismatch cleavage

- 5 \* Note: not useful for detection of promoter polymorphisms.

**Hybridisation Based:**

Solid phase hybridisation: Dot blots, MASDA, Reverse dot blots, Oligonucleotide arrays (DNA Chips).

Solution phase hybridisation: Taqman™ - US-5210015 & US-5487972 (Hoffmann-La

- 10 Roche), Molecular Beacons - Tyagi *et al* (1996), Nature Biotechnology, 14, 303; WO 95/13399 (Public Health Inst., New York).

**Extension Based:** ARMS™, ALEX™ - European Patent No. EP 332435 B1 (Zeneca Limited), COPS - Gibbs *et al* (1989), Nucleic Acids Research, 17, 2347.

**Incorporation Based:** Mini-sequencing, APEX

- 15 **Restriction Enzyme Based:** RFLP, Restriction site generating PCR

**Ligation Based:** OLA

**Other:** Invader assay

Table 2 - Signal Generation or Detection Systems

- 20 **Fluorescence:** FRET, Fluorescence quenching, Fluorescence polarisation - United Kingdom Patent No. 2228998 (Zeneca Limited)

**Other:** Chemiluminescence, Electrochemiluminescence, Raman, Radioactivity, Colorimetric, Hybridisation protection assay, Mass spectrometry.

- 25 Table 3 - Further Amplification Methods

SSR, NASBA, LCR, SDA, b-DNA

Preferred mutation detection techniques include ARMS™, ALEX™, COPS, Taqman, Molecular Beacons, RFLP, and restriction site based PCR and FRET techniques.

- 30 Particularly preferred methods include ARMS™ and RFLP based methods. ARMS™ is an especially preferred method.

In a further aspect, the diagnostic methods of the invention are used to assess the efficacy of therapeutic compounds in the treatment of PDH-mediated diseases such as diabetes, asthma, obesity, sepsis, and peripheral vascular disease.

Assays, for example reporter-based assays, may be devised to detect whether one or  
5 more of the above polymorphisms affect transcription levels and/or message stability.

Individuals who carry particular allelic variants of the PDH E1 $\beta$  gene may therefore exhibit differences in their ability to regulate protein biosynthesis under different physiological conditions and may display altered abilities to react to different diseases. In addition, differences in protein regulation arising as a result of allelic variation may have a  
10 direct effect on the response of an individual to drug therapy. The diagnostic methods of the invention may be useful both to predict the clinical response to such agents and to determine therapeutic dose.

In a further aspect, the diagnostic methods of the invention, are used to assess the predisposition of an individual to diseases mediated by PDH. This may be particularly  
15 relevant in the development of diabetes, asthma, obesity, sepsis, and peripheral vascular disease and other diseases which are mediated by PDH. The present invention may be used to recognise individuals who are particularly at risk from developing these conditions.

Low frequency polymorphisms may be particularly useful for haplotyping as described below. A haplotype is a set of alleles found at linked polymorphic sites (such as  
20 within a gene) on a single (paternal or maternal) chromosome. If recombination within the gene is random, there may be as many as  $2^n$  haplotypes, where 2 is the number of alleles at each polymorphic position and n is the number of polymorphic positions. One approach to identifying mutations or polymorphisms which are correlated with clinical response is to carry out an association study using all the haplotypes that can be identified in the population of  
25 interest. The frequency of each haplotype is limited by the frequency of its rarest allele, so that polymorphisms with low frequency alleles are particularly useful as markers of low frequency haplotypes. As particular mutations or polymorphisms associated with certain clinical features, such as adverse or abnormal events, are likely to be of low frequency within the population, low frequency polymorphisms may be particularly useful in identifying these  
30 mutations (for examples see: De Stefano V et al., *Ann Hum Genet* (1998) 62:481-90; and Keightley AM et al., *Blood* (1999) 93:4277-83).

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In a further aspect, the diagnostic methods of the invention are used in the development of new drug therapies which selectively target one or more allelic variants of the PDH E1 $\beta$  gene. Identification of a link between a particular allelic variant and predisposition to disease development or response to drug therapy may have a significant impact on the design of new drugs. Drugs may be designed to regulate the biological activity of variants implicated in the disease process whilst minimising effects on other variants.

In a further diagnostic aspect of the invention the presence or absence of variant nucleotides is detected by reference to the loss or gain of, optionally engineered, sites recognised by restriction enzymes.

- 10 According to another aspect of the present invention there is provided a nucleic acid comprising any one of the following polymorphisms:
- the nucleic acid of SEQ ID NO: 1 with G at position 457 as defined by the positions in SEQ ID NO: 1;
- the nucleic acid of SEQ ID NO: 1 with C at position 1191 as defined by the position in SEQ ID NO: 1;
- 15 the nucleic acid of SEQ ID NO: 1 with T at position 1198 as defined by the positions in SEQ ID NO: 1;
- the nucleic acid of SEQ ID NO: 1 with A at position 1342 as defined by the position in SEQ ID NO: 1;
- 20 or a complementary strand thereof or an antisense sequence thereto or a fragment thereof of at least 20 bases comprising at least one polymorphism.

Another aspect of the invention provides an isolated nucleic acid comprising the nucleic acid of SEQ ID NO: 1 with C at position 1191 as defined by the position in SEQ ID NO: 1; or a complementary strand thereof or an antisense sequence thereto or a fragment thereof of at least 20 bases comprising C at position 1191.

Fragments are at least 17 bases, more preferably at least 20 bases, more preferably at least 30 bases.

The scope of the invention does not extend to any nucleic acid as it is found in nature. A nucleic acid of the invention is preferably in isolated form, for example through being at least partially purified from any substance with which it occurs naturally (if any).

Novel sequence disclosed herein, may be used in another embodiment of the invention to regulate expression of the gene in cells by the use of antisense constructs. To enable methods of down-regulating expression of the gene of the present invention in mammalian cells, an example antisense expression construct can be readily constructed for

5 instance using the pREP10 vector (Invitrogen Corporation). Transcripts are expected to inhibit translation of the gene in cells transfected with this type of construct. Antisense transcripts are effective for inhibiting translation of the native gene transcript, and capable of inducing the effects (e.g., regulation of tissue physiology) herein described. Oligonucleotides which are complementary to and hybridisable with any portion of novel gene mRNA

10 disclosed herein are contemplated for therapeutic use. U.S. Patent No. 5,639,595, "Identification of Novel Drugs and Reagents", issued Jun. 17, 1997, wherein methods of identifying oligonucleotide sequences that display in vivo activity are thoroughly described, is herein incorporated by reference. Expression vectors containing random oligonucleotide sequences derived from previously known polynucleotides are transformed into cells. The

15 cells are then assayed for a phenotype resulting from the desired activity of the oligonucleotide. Once cells with the desired phenotype have been identified, the sequence of the oligonucleotide having the desired activity can be identified. Identification may be accomplished by recovering the vector or by polymerase chain reaction (PCR) amplification and sequencing the region containing the inserted nucleic acid material. Antisense molecules

20 can be synthesised for antisense therapy. These antisense molecules may be DNA, stable derivatives of DNA such as phosphorothioates or methylphosphonates, RNA, stable derivatives of RNA such as 2'-O-alkylRNA, or other oligonucleotide mimetics. U.S. Patent No. 5,652,355, "Hybrid Oligonucleotide Phosphorothioates", issued July 29, 1997, and U.S. Patent No. 5,652,356, "Inverted Chimeric and Hybrid Oligonucleotides", issued July 29,

25 1997, which describe the synthesis and effect of physiologically-stable antisense molecules, are incorporated by reference. Antisense molecules may be introduced into cells by microinjection, liposome encapsulation or by expression from vectors harboring the antisense sequence.

The invention further provides nucleotide primers which can detect the

30 polymorphisms of the invention.

According to another aspect of the present invention there is provided an allele specific primer capable of detecting a PDH E1 $\beta$  gene polymorphism at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1.

5 An allele specific primer is used, generally together with a constant primer, in an amplification reaction such as a PCR reaction, which provides the discrimination between alleles through selective amplification of one allele at a particular sequence position e.g. as used for ARMST<sup>TM</sup> assays. The allele specific primer is preferably 17- 50 nucleotides, more preferably about 17-35 nucleotides, more preferably about 17-30 nucleotides.

10 An allele specific primer preferably corresponds exactly with the allele to be detected but derivatives thereof are also contemplated wherein about 6-8 of the nucleotides at the 3' terminus correspond with the allele to be detected and wherein up to 10, such as up to 8, 6, 4, 2, or 1 of the remaining nucleotides may be varied without significantly affecting the properties of the primer.

15 Primers may be manufactured using any convenient method of synthesis. Examples of such methods may be found in standard textbooks, for example "Protocols for Oligonucleotides and Analogues; Synthesis and Properties," Methods in Molecular Biology Series; Volume 20; Ed. Sudhir Agrawal, Humana ISBN: 0-89603-247-7; 1993; 1<sup>st</sup> Edition. If required the primer(s) may be labelled to facilitate detection.

20 According to another aspect of the present invention there is provided an allele-specific oligonucleotide probe capable of detecting a PDH E1 $\beta$  gene polymorphism at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1.

The allele-specific oligonucleotide probe is preferably 17- 50 nucleotides, more  
25 preferably about 17-35 nucleotides, more preferably about 17-30 nucleotides.

The design of such probes will be apparent to the molecular biologist of ordinary skill. Such probes are of any convenient length such as up to 50 bases, up to 40 bases, more conveniently up to 30 bases in length, such as for example 8-25 or 8-15 bases in length. In general such probes will comprise base sequences entirely complementary to the  
30 corresponding wild type or variant locus in the gene. However, if required one or more mismatches may be introduced, provided that the discriminatory power of the oligonucleotide

probe is not unduly affected. The probes of the invention may carry one or more labels to facilitate detection.

According to another aspect of the present invention there is provided a diagnostic kit comprising an allele specific oligonucleotide probe of the invention and/or an allele-specific  
5 primer of the invention.

The diagnostic kits may comprise appropriate packaging and instructions for use in the methods of the invention. Such kits may further comprise appropriate buffer(s), nucleotides, and polymerase(s) such as thermostable polymerases, for example taq polymerase.

In another aspect of the invention, any polymorphism of this invention may be used as  
10 a genetic marker in a linkage study. This particularly applies to the polymorphisms at positions 457, 1191 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1 because of their relatively high frequency (see Examples below).

According to another aspect of the present invention there is provided a method of treating a human in need of treatment with a PDH drug in which the method comprises:

- 15 i) diagnosis of a polymorphism in the PDH E1 $\beta$  gene in the human, which diagnosis comprises determining the sequence of the nucleic acid at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1, and determining the status of the human by reference to polymorphism in the PDH E1 $\beta$  gene; and  
ii) administering an effective amount of a PDH drug.

20 Preferably determination of the status of the human is clinically useful. Examples of clinical usefulness include deciding which drug or drugs to administer and/or establishing the effective amount of the drug or drugs.

Drugs which increase the activity of PDH are of value in a number of disease conditions, including disease states associated with disorders of glucose utilisation such as  
25 diabetes and obesity, and associated with excessive production of lactate such as encountered in sepsis and other causes of lactic acidemia. Additionally drugs which increase the activity of PDH may be expected to have utility in diseases where supply of energy-rich substrates to tissues is limiting such as peripheral vascular disease, coronary failure and certain cardiac myopathies, muscle ataxia and weakness.

30 According to another aspect of the present invention there is provided use of a PDH drug in the preparation of a medicament for treating a PDH-mediated disease in a human

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diagnosed as having a polymorphism at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1.

According to another aspect of the present invention there is provided a pharmaceutical pack comprising a PDH drug and instructions for administration of the drug to humans diagnostically tested for a polymorphism at one or more of positions 457, 1191, 1198 and 1342 in the PDH E1 $\beta$  gene as defined by the positions in SEQ ID NO: 1.

According to another aspect of the present invention there is provided a computer readable medium comprising at least one novel polynucleotide sequence of the invention stored on the medium. The computer readable medium may be used, for example, in homology searching, mapping, haplotyping, genotyping or pharmacogenetic analysis or any other bioinformatic analysis. The reader is referred to Bioinformatics, A practical guide to the analysis of genes and proteins, Edited by A D Baxevanis & B F F Ouellette, John Wiley & Sons, 1998. Any computer readable medium may be used, for example, compact disk, tape, floppy disk, hard drive or computer chips.

The polynucleotide sequences of the invention, or parts thereof, particularly those relating to and identifying the polymorphisms identified herein represent a valuable information source, for example, to characterise individuals in terms of haplotype and other sub-groupings, such as investigation of susceptibility to treatment with particular drugs. These approaches are most easily facilitated by storing the sequence information in a computer readable medium and then using the information in standard bioinformatics programs or to search sequence databases using state of the art searching tools such as "GCC". Thus, the polynucleotide sequences of the invention are particularly useful as components in databases useful for sequence identity and other search analyses. As used herein, storage of the sequence information in a computer readable medium and use in sequence databases in relation to 'polynucleotide or polynucleotide sequence of the invention' covers any detectable chemical or physical characteristic of a polynucleotide of the invention that may be reduced to, converted into or stored in a tangible medium, such as a computer disk, preferably in a computer readable form. For example, chromatographic scan data or peak data, photographic scan or peak data, mass spectrographic data, sequence gel (or other) data.

The invention provides a computer readable medium having stored thereon one or more polynucleotide sequences of the invention. For example, a computer readable medium is provided comprising and having stored thereon a member selected from the group consisting of: a polynucleotide comprising the sequence of a polynucleotide of the invention, 5 a polynucleotide consisting of a polynucleotide of the invention, a polynucleotide which comprises part of a polynucleotide of the invention, which part includes at least one of the polymorphisms of the invention, a set of polynucleotide sequences wherein the set includes at least one polynucleotide sequence of the invention, a data set comprising or consisting of a polynucleotide sequence of the invention or a part thereof comprising at least one of the 10 polymorphisms identified herein.

A computer based method is also provided for performing sequence identification, said method comprising the steps of providing a polynucleotide sequence comprising a polymorphism of the invention in a computer readable medium; and comparing said polymorphism containing polynucleotide sequence to at least one other polynucleotide or 15 polypeptide sequence to identify identity (homology), i.e. screen for the presence of a polymorphism.

Another aspect of the invention provides the use of any one of the following in bioinformatic analysis:

- i) any polymorphism defined herein; or
- 20 ii) any haplotype defined herein.

Preferred uses comprise a bioinformatic analysis selected from homology searching, mapping, haplotyping, genotyping or pharmacogenetic analysis.

The invention will now be illustrated but not limited by reference to the following Examples. All temperatures are in degrees Celsius.

25 In the Examples below, unless otherwise stated, the following methodology and materials have been applied.

AMPLITAQ™ or AMPLITAQ GOLD™ available from Perkin-Elmer Cetus, are used as the source of thermostable DNA polymerase.

General molecular biology procedures can be followed from any of the methods 30 described in "Molecular Cloning - A Laboratory Manual" Second Edition, Sambrook, Fritsch and Maniatis (Cold Spring Harbor Laboratory, 1989).

Electropherograms were obtained in a standard manner: data was collected by ABI377 data collection software and the wave form generated by ABI Prism™ sequencing analysis (2.1.2).

#### Example 1

### 5 Identification of Polymorphisms

#### 1. Methods

##### c-DNA Preparation

RNA was prepared from lymphoblastoid cell lines from Caucasian donors using standard laboratory protocols (Chomczynski and Sacchi, Anal. Biochem. 162, 156-159, 1987) and used to generate first strand cDNA (Gubler and Hoffman, Gene 25, 263-269, 1983).

##### Template Preparation

Templates were prepared by PCR using the oligonucleotide primers and annealing temperatures set out below. The extension temperature was 72° and denaturation temperature 94°; each step was 1 minute. Generally 100 pg cDNA was used in each reaction and subjected to 40 cycles of PCR.

Fragment	Forward Oligo	Reverse Oligo	Annealing Temp	MgCl <sub>2</sub>
1-420	1-22	397-420	62°	2mM
299-791	299-320	770-791	63°	1mM
714-1194	714-736	1171-1194	62°	2mM
999-1410	999-1021	1388-1410	62°	2mM

All positions refer to the positions in SEQ ID NO: 1.

For dye-primer sequencing the forward primers were modified to include M13 forward sequence (ABI protocol P/N 402114, Applied Biosystems) at the 5' end of the oligonucleotides.

##### Dye Primer Sequencing

Dye-primer sequencing using M13 forward primer was as described in the ABI protocol P/N 402114 for the ABI Prism™ dye primer cycle sequencing core kit with

“AmpliTaq FS”™ DNA polymerase, modified in that the annealing temperature was 45° and DMSO was added to the cycle sequencing mix to a final concentration of 5%.

The extension reactions for each base were pooled, ethanol/sodium acetate precipitated, washed and resuspended in formamide loading buffer.

- 5 4.25% Acrylamide gels were run on an automated sequencer (ABI 377, Applied Biosystems).

## 2. Results

### Novel Polymorphisms

<i>Position</i>	<i>Published allele</i>	<i>Variant allele</i>	<i>amino acid change</i>	<i>RFLP</i>	<i>Allele Frequency</i>
457	A	G	silent Gly	+ Apa I	30/50
1191	A	C	3'UTR		32/54
1198	C	T	3'UTR		17/54
1342	C	A	3'UTR		32/54

All positions refer to the positions in SEQ ID NO: 1.

- 10 Frequency is the allele frequency of the variant allele in control subjects.

### *Three common haplotypes in the 3'UTR of PDH E1 $\beta$ :*

The 3'UTR of PDH E1 $\beta$  was sequenced from 27 individuals. 15 individuals were homozygous at positions 1191, 1198 and 1342 indicating three common haplotypes listed below.

- 15 (1) 1191C 1198C 1342A  
(2) 1191A 1198C 1342C  
(3) 1191C 1198T 1342A

The remaining 12 individuals were heterozygous at two or more of positions 1191, 1198 and 1342 but were all compatible with heterozygosity for the above haplotypes.

- 20 The inferred haplotype frequencies were:

- (1) 15/54  
(2) 22/54  
(3) 17/54